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Partially esterified or amidated lactic or glycolic acid oligomers - for preparing delayed release formes of pharmaceuticals e.g. calcitonin and bromocriptine

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Patent Family (16 patents, 8 countries)

Patent	Application					
Number	Kind	Date N	umber	Kind	Date 1	Jpdate
BE 898430	.A	19840612	BE 89843	80 z	4 198312	212 198426 B
DE 3345314	\mathbf{A}	19840705	DE 3345	314	A 1983.	1215 198428 E
FR 2537980	A	19840622	FR 1983	19976	A 1983	31212 198430 E
GB 2135320	Α	19840830	GB 1983	33289	A 198	31214 198435 E
JP 59130252	A	19840726	JP 19832	38721	A 1983	1216 198436 E
CH 651475	A	19850930	CH 1982	7372	A 1982	1217 198542 E
CH 19827373 A 19821217						
GB 2165454	\mathbf{A}	19860416	GB 1985	27420	A 198	51107 198616 E
GB 2165849	\mathbf{A}	19860423	GB 1985	527421	A 198	51107 198617 E
GB 2166652	A	19860514			19862	20 E
GB 2165454	В	19870311			19871	0 E
GB 2166652	В	19870311			19871	0 E
GB 2135320	В	19870429	•		19871	7 E
CH 660488	A	19870430	CH 1982	7372	A 1982	1217 198720 E
CH 19827373 A 19821217						
CH 663718	Α	19880115	CH 1982	7372	A 1982	1217 198807 E
CH 19827373 A 19821217						
US 4801739	A	19890131	US 1983	562470	A 198	31216 198907 E
IT 1197759	В	19881206			199113	8 E

Priority Applications (no., kind, date): CH 19827372 A 19821217; CH 19827373 A 19821217

Alerting Abstract BE A

The oligomers have a mol. wt. of 500-10,000. The free carboxy gp. of the oligomer is at least partially in the form of an amide gp. with an aminoacid or in the form of an ester gp. with a sterol. The prods. pref. have an acid index of 1.5 or less.

Used for prepn, of delayed release forms of pharmacologically active cpds. The oligomers allow a sufficiently long release time for the active cpds, e.g. several weeks, and are appropriate for use with a wide variety of active cpds., such as those soluble in water. The oligomers are easy to manipulate, are less adhesive than polymers having free acid gps, and may be administered in the form of microcapsules (e.g. 10-60 microns) by a

syringe, e.g. intramuscularly or sub-cutaneously.

Equivalent Alerting Abstract US A

Modified oligomer comprises units of lactic acid and/or glycolic acid. The oligomer moiety has a mol, wt. of 500-10,000, pref. 750-5000, in which the terminal carboxy gp. is in the form of an amide or an alpha amino acid, pref. an alpha amino acid amide, esp. phenylaniline.

Pref. the oligomer moiety contains lactic acid units.

USE - New oligomer is useful in the prod. of depot forms of pharmacologically active agents including salmon calcitonin and bromocriptine.

USE - (7pp)